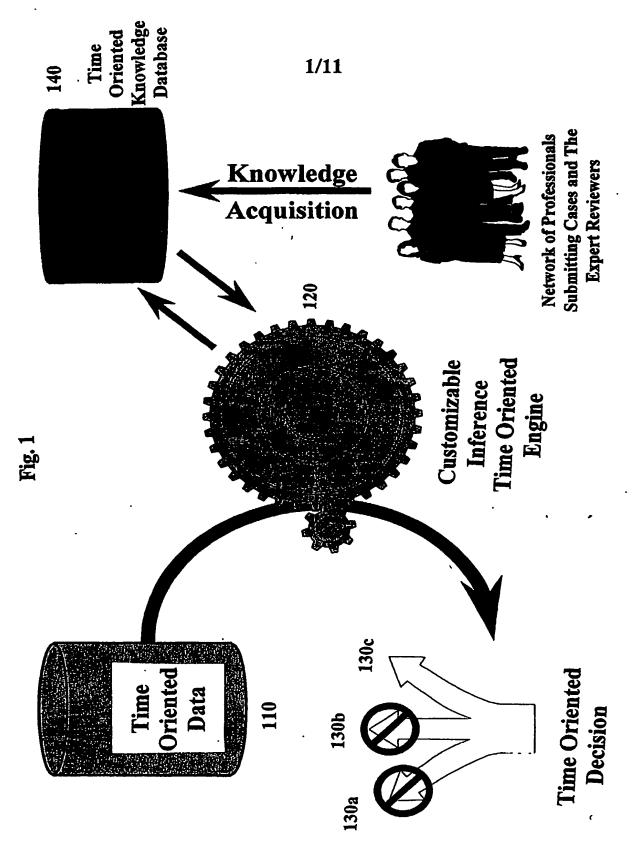
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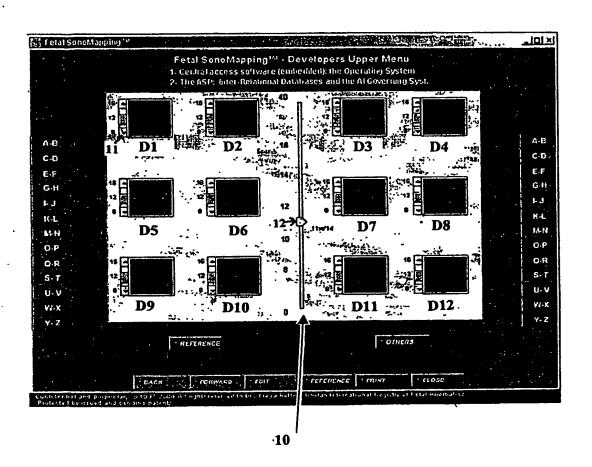


Fig. 2

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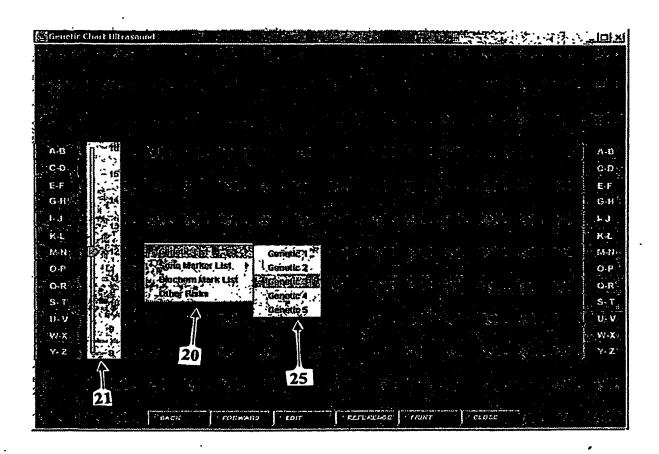


Fig. 2A

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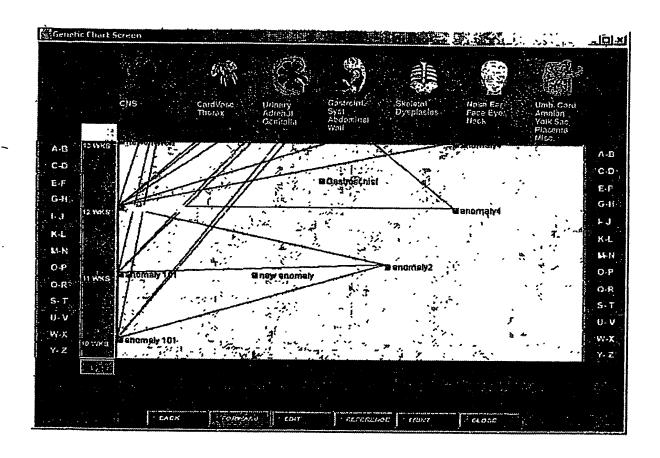


Fig. 2B

Algo based new associated sigr											3/	I										
Algo Based Alternative Syndromes by Probability with Polydactyry at 10 wks, After Earliest seen&Classif Exclusion of Meckal_Gruber Syndrome	Bardet-Bedi syndrame	Ellis-Van Crevold syndrome	Short rib polydectyly syndrome Type 1	Palister-Killan Syndrome	46, XY, -5, +den(5)	APERT SYNDROME	BECKWITH-WIEDEMANN SYNDROME; BWS	HOLT-ORAM SYNDROME; HOS	ACHONDROGENESIS, TYPE II; ACG2	FRASER SYNDROME	RUTLEDGE LETHAL MULTIPLE CONGENITAL ANOMALY SYNDROME	SMITH-LEMU-OPITZ SYNDROME; SLOS	SIMPSON-GÖLABLBEHMEL SYNDROME, TYPE 1; SGBS1	ACROCALLOSAL SYNDROME; ACLS	ASPHYXIATING THORACIC DYSTROPHY; ATD	JOUBERT SYNDROME 1; JBTS1	HYDROLETHALUS SYNDROME	SPLIT-HAND/FOOT MALFORMATION 3; SHFM3	ECTODERMAL DYSPLASIA SYNDROME WITH DISTINCTIVE FACIAL APPEARANCE AND PREAXIAL POLYDACTYLY OF FEET	VATER ASSOCIATION	DANDY-WALKER MALFORMATION WITH POSTAXIAL POLYDACTYLY	JOUBERT SYNDROME WITH BILATERAL CHORIORETINAL COLOBOMA
Exrilest seen&Classif			1 SMS		11 whos i	12wks 1	towics I	15wks	flwts I	11wks			10 WKS (I	11wks 1	11wits I	15wks				10wfs 1	10wfcs III	
Algo Based Associated Signs		Sloping forehead	Micrognathia .	Potter-like facies	Low-set ears	Microphthalmia	Hypotelorism	Hypertetorism	Cleft palate	cleft (b	Macrostomia	Short neck	Increased NT	Septal defects	Coarctation of sorts	Pulmonary hypoplasia	Splenamegaly	Asplenia	Accessory spleen	Single umbilical artery 10wts	Omphelocele	Intestinal maintation
Algo Based Most Common Algo Based Syndrome Associated	Meckel Gruber Syndrome																	,				
Earliest Seen	10wks																					
Earth	Polydactyly (101)																					

Fig. 2(

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		Imperforate anus		ORAL-FACIAL-DIGITAL SYNDROME, TYPE IV
		Small centralia	16wts (small penis)	MIDLINE MALFORMATIONS, MULTIPLE, WITH LIMB ABNORMALITIES AND HYPOPITUITARISM
		allal	15wks	HOLOPROSENCEPHALY 1, ALOBAR; HPE1
		Cryptorchidism		ROBINOW-SORAUF SYNDROME
		Polycystic kidneys	III STAOS	THANATOPHORIC DYSPLASIA; TD
			12wts 1	RUBINSTEIN SYNDROME
		Duplicated uniters		KLIPPEL-TRENAUNAY-WEBER SYNDROME
		Hypoplastic bladder		POLYCYSTIC KIDNEYS
			18w/cs	
			9wits 1	30
		,	15m/cs	
		Tellpes	11wks III	
		Polydactyly (101)	10wks 1	
		Amoid-Chiari metformation	1 dw/cs	
		Occipital		
		encephalocele	10wfcs 1	
		Hydrocephatus	9wtcs III	
		Dandy-Walker matformation	11wtes III	
		Cerebral hypoplasia		
			14wts	
		Anencephaly	9wice 1	
	`	Absence of corpus cellosum		
		Adrenal hypoplasia		
		Oligohydramnios	9wks III	
		Placental enlargement		

**Fig. 2I** 

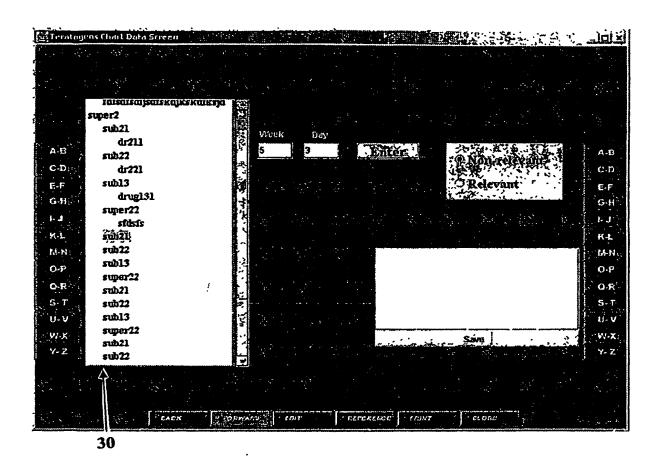


Fig. 3A

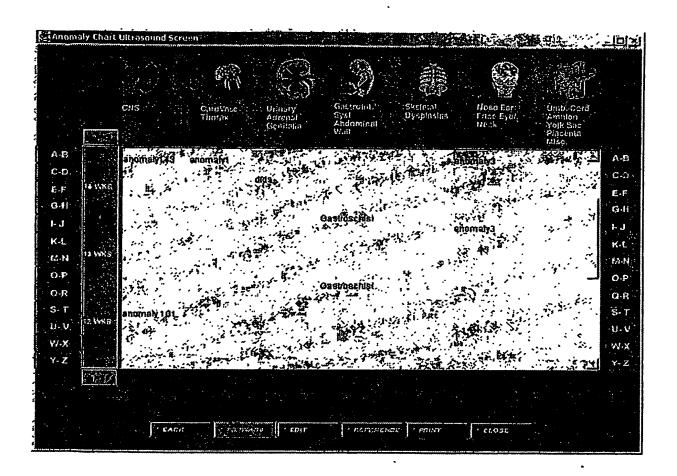


Fig. 3B

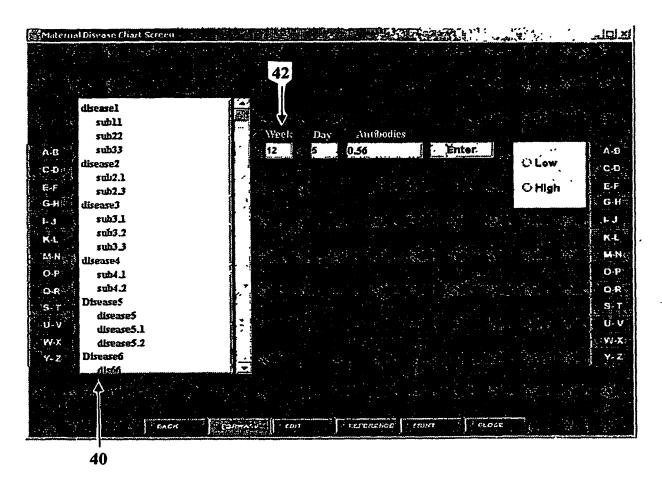


Fig. 4A

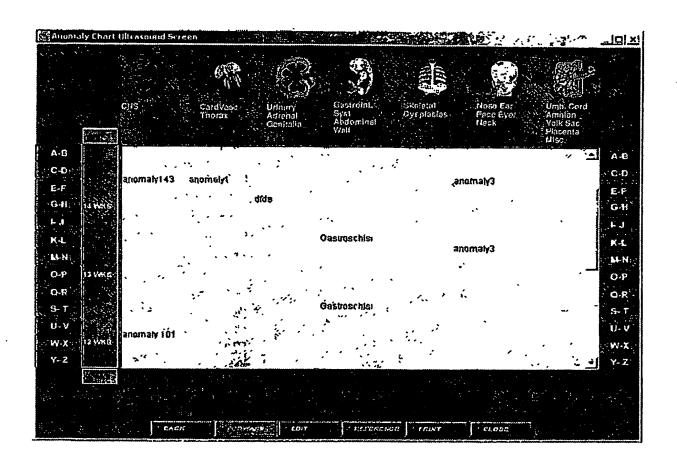


Fig. 4B

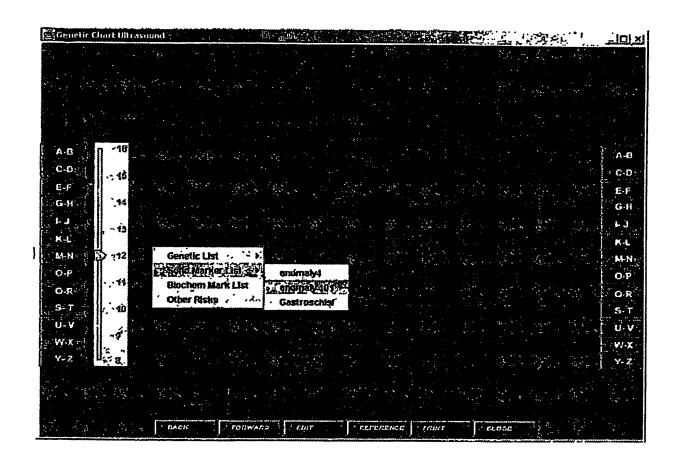


Fig. 4C

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